Strontium-90 was only detected at WEE, depth 0-2cm, slightly above the MDC (Table 4.13). However, ⁶⁰Co was not detected in any of the soil samples. There was nonsignificant difference in ⁶⁰Co concentrations between years (p = 0.421).

Soil samples collected from one location (WEE) were divided into two parts and analyzed separately (Table 4.14). Uranium-234, ²³⁸U, ⁴⁰K, and ¹³⁷Cs were compared between the duplicates. Other radionuclides of interest had insufficient detections to allow a reasonable comparison. The RER was greater than one for ²³⁴U and ²³⁸U in all samples. However, it was less than one in one for ⁴⁰K analyses and ¹³⁷Cs analyses. A paired t-test indicated no significant difference between ²³⁴U duplicates (p = 0.624) and between ²³⁸U duplicates (p = 0.666). This circumstance indicates a lack of precision in these analyses, primarily due to the non-homogeneous distribution of radionuclides in soils. Because of small-scale differences in topography, soil type and structure, soil moisture, and other microenvironmental conditions, radionuclides are rarely homogeneously distributed in soils, and good agreement between duplicate samples is difficult to achieve. However, all the measurements were low, within the range of natural concentrations, and did not differ in time or space in such a way as to suggest WIPP-related contamination of the environment.

Table 4.14 - Results of Duplicate Soil Sample Analysis. Units are Bq/g. See Appendix B for the sampling locations.

	The second secon											
	Depth	[RN] ^a	2×TPU ^b	MDC°	RER ^d	[RN]	2×TPU ^a	MDC ^b	RER ^c			
Location	(cm)		²³⁴ U				²³⁸ U					
WEE	0-2	5.66×10 ⁻³	1.35×10 ⁻³	1.02×10 ⁻⁴	2.94	6.85×10 ⁻³	1.57×10 ⁻³	1.01×10 ⁻⁴	2.26			
WEE De	0-2	1.50×10 ⁻²	2.88×10 ⁻³	8.92×10 ⁻⁴		1.39×10 ⁻²	2.70×10 ⁻³	8.88×10 ⁻⁵				
WEE	2-5	6.73×10 ⁻³	1.38×10 ⁻³	7.81×10 ⁻⁵	2.63	7.33×10 ⁻³	1.48×10 ⁻³	7.77×10 ⁻⁵	2.81			
WEE De	2-5	1.57×10 ⁻²	3.12×10 ⁻³	9.99×10 ⁻⁵		1.81×10 ⁻²	3.53×10 ⁻³	9.92×10 ⁻⁵				
WEE	5-10	1.84×10 ⁻²	3.68×10 ⁻³	2.83×10 ⁻⁴	1.94	2.00×10 ⁻²	3.96×10 ⁻³	1.04×10 ⁻⁴	1.90			
WEE De	5-10	1.01×10 ⁻²	2.20×10 ⁻³	3.00×10 ⁻⁴		1.12×10 ⁻²	2.39×10 ⁻³	3.77×10 ⁻⁴				
			⁴⁰ K				¹³⁷ Cs					
WEE	0-2	2.25×10 ⁻¹	3.07×10 ⁻²	7.36×10 ⁻³	0.40	2.21×10 ⁻³	5.48×10 ⁻⁴	6.55×10 ⁻⁴	0.25			
WEE De	0-2	2.43×10 ⁻¹	3.30×10 ⁻²	6.29×10 ⁻³		2.40×10 ⁻³	5.22×10 ⁻⁴	5.55×10 ⁻⁴				
WEE	2-5	2.62×10 ⁻¹	3.55×10 ⁻²	1.32×10 ⁻²	0.41	2.49×10 ⁻³	6.22×10 ⁻⁴	7.62×10 ⁻⁴	0.18			
WEE De	2-5	2.42×10 ⁻¹	3.28×10 ⁻²	6.36×10 ⁻³		2.35×10 ⁻³	4.74×10 ⁻⁴	4.55×10 ⁻⁴				
WEE	5-10	2.62×10 ⁻¹	3.55×10 ⁻²	1.32×10 ⁻²	0.22	2.49×10 ⁻³	6.22×10 ⁻⁴	7.62×10 ⁻⁴	0.50			
WEE De	5-10	2.51×10 ⁻¹	3.42×10 ⁻²	1.34×10 ⁻²		2.12×10 ⁻³	4.07×10 ⁻⁴	5.14×10 ⁻⁴				

a [RN] = Radionuclide concentration

4.7 **Sediments**

4.7.1 Sample Collection

Sediment samples were collected from 12 locations around the WIPP site, mostly from the same water bodies from which the surface water samples were collected (Figure 4.7, see Appendix B for location codes). The samples were collected in 1 l

^b Total propagated uncertainty

^c Minimum detectable concentration

^d Relative Error Ratio

^e Duplicate

plastic containers from the top 15 cm (6 in) of the sediments of the water bodies and shipped to the laboratory for the determination of individual radionuclides.

4.7.2 Sample Preparation

Sediment samples were dried at 110°C (230°F) for several hours and homogenized by grinding to smaller particle sizes. A 0.75-g (0.04-oz) aliquot was dissolved by heating it with a mixture of nitric, hydrochloric, and hydrofluoric acids. The residue was heated with nitric and boric acids to remove hydrofluoric acid quantitatively. Finally, the residue was dissolved in hydrochloric acid for the determination of individual radionuclides.

4.7.3 Determination of Individual Radionuclides

About 100 g (4 oz) of dried and homogenized sediment samples were counted by gamma-spectrometry for the determinations of ⁴⁰K, ⁶⁰Co, and ¹³⁷Cs. Strontium-90 was determined from an aliquot of dissolved sediment samples by chemical separation and beta counting. Uranium, plutonium, and americium were determined by alpha spectrometry after chemical separations, micro-precipitating, and filtering onto micro filter papers.

4.7.4 Results and Discussions

Uranium-234, 235 U, and 238 U were detected in every sediment sample in (Table 4.15). The concentration of 234 U ranged from $1.73\times10^{-2}\pm3.10\times10^{-3}$ Bq/g ($4.68\times10^{-1}\pm8.38\times10^{-2}$ pCi/g) at LST to $4.96\times10^{-2}\pm8.33\times10^{-3}$ Bq/g ($1.34\times10^{0}\pm2.25\times10^{-1}$ pCi/g) at CBD. The concentration of 235 U ranged from $9.32\times10^{-4}\pm3.60\times10^{-4}$ Bq/g ($2.52\times10^{-2}\pm9.73\times10^{-3}$ pCi/g) at RED to $2.12\times10^{-3}\pm6.25\times10^{-4}$ Bq/g ($5.73\times10^{-2}\pm1.69\times10^{-2}$ pCi/g) at CBD. The concentration of 238 U was lowest at LST ($1.84\times10^{-2}\pm3.27\times10^{-3}$ Bq/g [$4.97\times10^{-1}\pm8.84\times10^{-2}$ pCi/g]) and highest at CBD ($3.35\times10^{-2}\pm5.74\times10^{-4}$ Bq/g [$9.05\times10^{-1}\pm1.55\times10^{-2}$ pCi/g]). As expected, the 235 U concentration was much lower than the concentrations of 234 U and 238 U. There was not a significant difference between sampling years (ANOVA 234 Up = 0.526, 235 Up = 0.399, 238 U p = 0.315).

Plutonium-238 was not detected in any sediment sample in 2001 (Table 4.16). Americium-241 was measured above the MDC at BRA. Plutonium-239+240 concentration was greater than the MDC at BRA, IDN, and Poker Trap [PKT]. These samples were processed several times in the laboratory due to interference in the spectra. The samples showed concentration barely above the MDC for two out of three analysis. The apparent lack of homogeneity of the samples caused the different results. Sample location BRA indicated ²⁴¹Am concentration above the MDC. This sample was also reprocessed for spectral interference. The radionuclide concentrations in Table 4.16 are the reprocessed results.

Cesium-137 was detected in all the sediment samples, ranging from 4.14×10^{-4} ± 1.60×10^{-4} Bq/g (1.12×10^{-2} ± 4.32×10^{-3} pCi/g) at UPR to 4.59×10^{-2} ± 5.77×10^{-3} Bq/g (1.24×10^{0} ± 1.56×10^{-1} pCi/g) at BRA (Table 4.17). Cesium-137 did not differ statistically between sampling years 2000 and 2001 (ANOVA p = 0.397).

Strontium-90 was not detected in any sediment samples and Cobalt-60 was detected in one sediment sample (BRA). None of these radionuclides had sufficient detections to justify statistical comparisons between locations or years.

Potassium-40 was detected, as expected, in all sediment samples (Table 4.17). Its lowest concentration was found at CBD $(2.49\times10^{-1}\pm3.25\times10^{-2}\text{ Bq/g} [6.73\times10^{0}\pm8.78\times10^{-1}\text{ pCi/g}])$ and its highest concentration was found at TUT($1.04\times10^{0}\pm1.34\times10^{-1}$ Bq/g $[2.81\times10^{1}\pm3.62\times10^{-1}\text{ pCi/g}])$. Potassium-40 did not vary significantly between years (ANOVA, p = 0.977). Overall, the concentrations measured in 2001 were similar to the average concentration of 40 K found in soils throughout the United States $(4.00\times10^{-1}\text{ Bq/g} [1.08\times10^{1}\text{ pCi/g}]; NCRP, 1994)$.

Duplicate analyses were performed for all the radionuclides in sediment samples IDN and TUT (Table 4.18). The RER was less than one for 241 Am and all uranium isotopes, indicating acceptable correspondence between the original and the duplicate samples. For 137 Cs and 40 K, it was greater than one for location TUT. However, a t-test indicated no significant difference between any of these duplicate measurements for 40 K (p = 0.613) and 137 Cs (p = 0.723).

Table 4.15 - Uranium Concentrations (Bq/g) in Sediment Near the WIPP Site. See Appendix B for the sampling locations.

	[RN] ^a	2 × TPU ^b	MDC ^c	[RN]	2 × TPU	MDC	[RN]	2 × TPU	MDC
Location		²³⁴ U			²³⁵ U			²³⁸ U	
BRA	2.82×10 ⁻²	4.81×10 ⁻³	6.25×10 ⁻⁵	1.51×10 ⁻³	4.81×10 ⁻⁴	7.73×10 ⁻⁵	2.61×10 ⁻²	4.48×10 ⁻³	6.25×10 ⁻⁵
BHT	2.15×10 ⁻²	3.67×10 ⁻³	5.99×10 ⁻⁵	1.25×10 ⁻³	4.18×10 ⁻⁴	7.40×10 ⁻⁵	2.32×10 ⁻²	3.96×10 ⁻³	1.62×10 ⁻⁴
CBD	4.96×10 ⁻²	8.33×10 ⁻³	7.07×10 ⁻⁵	2.12×10 ⁻³	6.25×10 ⁻⁴	8.70×10 ⁻⁵	3.35×10 ⁻²	5.74×10 ⁻³	1.91×10 ⁻⁴
HIL	1.92×10 ⁻²	3.51×10 ⁻³	7.18×10 ⁻⁵	1.50×10 ⁻³	5.11×10 ⁻⁴	8.84×10 ⁻⁵	2.09×10 ⁻²	3.77×10 ⁻³	1.94×10 ⁻⁴
IDN	2.33×10 ⁻²	3.89×10 ⁻³	5.40×10 ⁻⁵	1.23×10 ⁻³	4.03×10 ⁻⁴	1.81×10 ⁻⁴	2.30×10 ⁻²	3.85×10 ⁻³	5.37×10 ⁻⁵
LST	1.73×10 ⁻²	3.10×10 ⁻³	2.21×10 ⁻⁴	1.15×10 ⁻³	4.11×10 ⁻⁴	7.96×10 ⁻⁵	1.84×10 ⁻²	3.27×10 ⁻³	6.44×10 ⁻⁵
NOY	2.32×10 ⁻²	3.89×10 ⁻³	5.81×10 ⁻⁵	1.21×10 ⁻³	4.07×10 ⁻⁴	7.14×10 ⁻⁵	2.43×10 ⁻²	4.07×10 ⁻³	5.77×10 ⁻⁵
PCN	3.56×10 ⁻²	6.29×10 ⁻³	2.00×10 ⁻⁴	1.91×10 ⁻³	5.99×10 ⁻⁴	9.07×10 ⁻⁵	3.20×10 ⁻²	5.66×10 ⁻³	7.33×10 ⁻⁵
PKT	2.55×10 ⁻²	4.18×10 ⁻³	1.53×10 ⁻⁴	1.94×10 ⁻³	5.37×10 ⁻⁴	6.92×10 ⁻⁵	2.70×10 ⁻²	4.40×10 ⁻³	5.59×10 ⁻⁵
RED	1.94×10 ⁻²	3.34×10 ⁻³	1.74×10 ⁻⁴	9.32×10 ⁻⁴	3.60×10 ⁻⁴	7.88×10 ⁻⁵	1.92×10 ⁻²	3.30×10 ⁻⁴	6.36×10 ⁻⁵
TUT	2.32×10 ⁻²	3.89×10 ⁻³	6.62×10 ⁻⁵	1.63×10 ⁻³	5.07×10 ⁻⁴	8.18×10 ⁻⁵	2.53×10 ⁻²	4.18×10 ⁻³	6.59×10 ⁻⁵
UPR	2.60×10 ⁻²	4.48×10 ⁻³	6.40×10 ⁻⁵	1.31×10 ⁻³	4.51×10 ⁻⁴	2.15×10 ⁻⁴	2.69×10 ⁻²	4.63×10 ⁻³	6.36×10 ⁻⁵

^a [RN] = Radionuclide concentration

^b Total propagated uncertainty

^c Minimum detectable concentration

Table 4.16 - Americium and Plutonium Concentrations (Bq/g) in Sediment Near the WIPP Site. See Appendix B for the sampling locations.

-	[RN] ^a	2 × TPU ^b	MDC°	[RN]	2 × TPU	MDC	[RN]	2 × TPU	MDC
Location		²⁴¹ Am			²³⁸ Pu			²³⁹⁺²⁴⁰ Pu	
BRA	7.10×10 ⁻⁴	4.03×10 ⁻⁴	1.37×10 ⁻⁴	1.23×10 ⁻⁴	1.25×10 ⁻⁴	8.33×10 ⁻⁵	2.46×10 ⁻³	6.85×10 ⁻⁴	8.33×10 ⁻⁵
BHT	2.04×10 ⁻⁴	2.18×10 ⁻⁴	3.16×10 ⁻⁴	0.00×10 ⁰	0.00×10 ⁰	3.03×10 ⁻⁴	1.64×10 ⁻⁴	1.67×10 ⁻⁴	1.11×10 ⁻⁴
CBD	3.06×10 ⁻⁵	1.37×10 ⁻⁴	2.84×10 ⁻⁴	1.02×10 ⁻⁴	1.04×10 ⁻⁴	6.96×10 ⁻⁵	7.70×10 ⁻⁵	8.95×10 ⁻⁵	6.96×10 ⁻⁵
HIL	1.51×10 ⁻⁴	1.62×10 ⁻⁴	2.22×10 ⁻⁴	-8.14×10 ⁻⁵	1.16×10 ⁻⁴	3.77×10 ⁻⁴	1.62×10 ⁻⁴	1.65×10 ⁻⁴	1.10×10 ⁻⁴
IDN	0.00×10^{0}	0.00×10°	2.30×10 ⁻⁴	-3.96×10 ⁻⁵	1.38×10 ⁻⁴	3.68×10 ⁻⁴	7.92×10 ⁻⁴	3.85×10 ⁻⁴	1.07×10 ⁻⁴
LST	1.10×10 ⁻⁴	1.57×10 ⁻⁴	2.55×10 ⁻⁴	6.85×10 ⁻⁵	1.37×10 ⁻⁴	2.51×10 ⁻⁴	1.36×10 ⁻⁴	1.38×10 ⁻⁴	9.25×10 ⁻⁵
NOY	1.43×10 ⁻⁴	1.74×10 ⁻⁴	2.66×10 ⁻⁴	3.89×10 ⁻⁵	1.35×10 ⁻⁴	2.87×10 ⁻⁴	1.95×10 ⁻⁴	2.08×10 ⁻⁴	2.87×10 ⁻⁴
PCN	0.00×10 ⁰	0.00×10°	2.19×10 ⁻⁴	0.00×10°	0.00×10 ⁰	5.03×10 ⁻⁴	0.00×10°	0.00×10 ⁰	1.47×10 ⁻⁴
PKT	7.73×10 ⁻⁵	1.10×10 ⁻⁴	1.05×10 ⁻⁴	7.88×10 ⁻⁵	1.58×10 ⁻⁴	2.90×10 ⁻⁴	7.07×10 ⁻⁴	3.74×10 ⁻⁴	2.90×10 ⁻⁴
RED	8.25×10 ⁻⁵	1.24×10 ⁻⁴	2.02×10 ⁻⁴	7.77×10 ⁻⁵	1.16×10 ⁻⁴	1.90×10 ⁻⁴	7.73×10 ⁻⁵	9.03×10 ⁻⁵	6.99×10 ⁻⁵
TUT	3.05×10 ⁻⁵	1.06×10 ⁻⁴	2.25×10 ⁻⁴	-2.51×10 ⁻⁵	5.03×10 ⁻⁵	1.85×10 ⁻⁴	1.25×10 ⁻⁴	1.34×10 ⁻⁴	1.85×10 ⁻⁴
UPR	0.00×10 ⁰	0.00×10°	7.10×10 ⁻⁵	-5.22×10 ⁻⁵	1.05×10 ⁻⁴	2.81×10 ⁻⁴	7.81×10 ⁻⁵	9.10×10 ⁻⁵	7.07×10 ⁻⁵

^a [RN] = Radionuclide concentration ^b Total propagated uncertainty ^c Minimum detectable concentration

Table 4.17 - Selected Radionuclide Concentrations (Bq/g) in Sediment Near the WIPP Site. See Appendix B for the sampling locations.

	[RN] ^a	2 × TPU ^b	MDC°	[RN]	2 × TPU	MDC			
Location		¹³⁷ Cs			⁶⁰ Co				
BRA	4.59×10 ⁻²	5.77×10 ⁻³	4.22×10 ⁻⁴	6.85×10 ⁻⁴	5.85×10 ⁻⁴	6.73×10 ⁻⁴			
BHT	6.77×10 ⁻³	9.03×10 ⁻⁴	3.58×10 ⁻⁴	4.14×10 ⁻⁵	5.88×10 ⁻⁴	6.59×10 ⁻⁴			
CBD	2.51×10 ⁻³	4.74×10 ⁻⁴	4.77×10 ⁻⁴	2.60×10 ⁻⁴	5.07×10 ⁻⁴	5.85×10 ⁻⁴			
HIL	7.99×10 ⁻³	1.10×10 ⁻³	5.14×10 ⁻⁴	8.40×10 ⁻⁵	6.25×10 ⁻⁴	7.10×10 ⁻⁴			
IDN	5.85×10 ⁻³	7.84×10 ⁻⁴	3.36×10 ⁻⁴	-1.18×10 ⁻⁴	2.50×10 ⁻⁴	2.74×10 ⁻⁴			
LST	1.73×10 ⁻³	4.14×10 ⁻⁴	5.11×10 ⁻⁴	5.40×10 ⁻⁴	5.07×10 ⁻⁴	5.88×10 ⁻⁴			
NOY	8.92×10 ⁻⁴	2.64×10 ⁻⁴	3.47×10 ⁻⁴	2.36×10 ⁻⁴	4.85×10 ⁻⁴	5.55×10 ⁻⁴			
PCN	7.62×10 ⁻⁴	1.69×10 ⁻⁴	2.55×10 ⁻⁴	1.55×10⁻⁴	4.51×10 ⁻⁴	5.18×10 ⁻⁴			
PKT	1.61×10 ⁻²	2.03×10 ⁻³	4.26×10 ⁻⁴	3.42×10 ⁻⁴	7.07×10 ⁻⁴	7.99×10 ⁻⁴			
RED	4.07×10 ⁻³	5.77×10 ⁻⁴	3.70×10 ⁻⁴	5.00×10 ⁻⁴	5.48×10 ⁻⁴	6.33×10 ⁻⁴			
TUT	3.81×10 ⁻³	5.62×10 ⁻⁴	4.44×10 ⁻⁴	3.81×10 ⁻⁴	7.29×10 ⁻⁴	8.25×10 ⁻⁴			
UPR	4.14×10 ⁻⁴	1.60×10 ⁻⁴	3.57×10 ⁻⁴	5.37×10 ⁻⁴	5.33×10 ⁻⁴	6.18×10 ⁻⁴			
		90Sr		<u>⁴⁰</u> K					
BRA	1.70×10 ⁻²	1.55×10 ⁻²	2.49×10 ⁻²	5.51×10 ⁻¹	7.10×10 ⁻²	6.51×10 ⁻³			
BHT	1.45×10 ⁻²	1.16×10 ⁻²	1.80×10 ⁻²	6.66×10 ⁻¹	8.55×10 ⁻²	5.96×10 ⁻³			
CBD	1.79×10 ⁻³	6.59×10 ⁻³	1.11×10 ⁻²	2.49×10 ⁻¹	3.25×10 ⁻²	5.59×10 ⁻³			
HIL	6.85×10 ⁻³	7.59×10 ⁻³	1.20×10 ⁻²	8.66×10 ⁻¹	1.11×10 ⁻¹	6.55×10 ⁻³			
IDN	-5.48×10 ⁻⁴	8.03×10 ⁻³	1.35×10 ⁻²	4.81×10 ⁻¹	6.18×10 ⁻²	6.07×10 ⁻³			
LST	-3.07×10 ⁻³	6.44×10 ⁻³	1.10×10 ⁻²	3.89×10 ⁻¹	5.03×10 ⁻²	6.03×10 ⁻³			
NOY	-7.22×10 ⁻⁴	6.07×10 ⁻³	1.02×10 ⁻²	4.77×10 ⁻¹	6.18×10 ⁻²	6.25×10 ⁻³			
PCN	1.54×10 ⁻³	6.66×10 ⁻³	1.10×10 ⁻²	3.27×10 ⁻¹	4.26×10 ⁻²	5.33×10 ⁻³			
PKT	2.16×10 ⁻³	7.07×10 ⁻³	1.16×10 ⁻²	8.70×10 ⁻¹	1.10×10 ⁻¹	6.99×10 ⁻³			
RED	1.32×10 ⁻³	6.92×10 ⁻³	1.15×10 ⁻²	4.85×10 ⁻¹	6.25×10 ⁻²	6.18×10 ⁻³			
TUT	2.52×10 ⁻³	7.14×10 ⁻³	1.20×10 ⁻²	1.04×10 ⁰	1.34×10 ⁻¹	7.77×10 ⁻³			
UPR	-2.64×10 ⁻⁴	6.99×10 ⁻³	1.20×10 ⁻²	4.81×10 ⁻¹	6.22×10 ⁻²	6.07×10 ⁻³			

a [RN] = Radionuclide concentration
b Total propagated uncertainty
c Minimum detectable concentration

Table 4.18 - Results of Duplicate Sediment Sample Analysis. Units are Bq/g. See Appendix B for the sampling locations.

	[RN] ^a	2×TPU ^b	MDC°	RER⁴	[RN]	2×TPU ^a	MDC ^b	RER°
Location		²⁴¹ A r	n	¹³⁷ Cs				
IDN	0.00×10°	0.00×10°	2.30×10 ⁻⁴	0.811	5.85×10 ⁻³	7.84×10 ⁻⁴	3.36×10 ⁻⁴	0.324
IDN Dup.	1.20×10 ⁻⁴	1.48×10 ⁻⁴	2.20×10 ⁻⁴		6.22×10 ⁻³	8.29×10 ⁻⁴	3.24×10 ⁻⁴	
TUT	3.05×10 ⁻⁵	1.06×10 ⁻⁴	2.25×10 ⁻⁴	0.492	3.81×10 ⁻³	5.62×10 ⁻⁴	4.44×10 ⁻⁴	1.419
TUT Dup.	6.59×10 ⁻⁵	9.40×10 ⁻⁵	8.92×10 ⁻⁵		2.80×10 ⁻³	4.37×10 ⁻⁴	4.22×10 ⁻⁴	
		⁴⁰ K		²³⁴ U				
IDN	4.81×10 ⁻¹	6.18×10 ⁻²	6.07×10 ⁻³	0.520	2.33×10 ⁻²	3.89×10 ⁻³	5.40×10 ⁻⁵	0.261
IDN Dup.	5.29×10 ⁻¹	6.85×10 ⁻²	5.18×10 ⁻³		2.19×10 ⁻²	3.69×10 ⁻³	5.55×10 ⁻⁵	
TUT	1.04×10°	1.34×10 ⁻¹	7.77×10 ⁻³	1.604	2.32×10 ⁻²	3.89×10 ⁻³	6.62×10 ⁻⁵	0.247
TUT Dup.	7.73×10 ⁻¹	9.88×10 ⁻²	6.66×10 ⁻³		2.46×10 ⁻²	4.11×10 ⁻³	1.67×10 ⁻⁴	
		²³⁵ U				²³⁸ U		
IDN	1.23×10 ⁻³	4.03×10 ⁻⁴	1.81×10 ⁻⁴	0.252	2.30×10 ⁻²	3.85×10 ⁻³	5.37×10 ⁻⁵	0.161
IDN Dup.	1.09×10 ⁻³	3.81×10 ⁻⁴	1.86×10 ⁻⁴		2.39×10 ⁻²	4.03×10 ⁻³	5.55×10 ⁻⁵	
TUT	1.63×10 ⁻³	5.07×10 ⁻⁴	8.18×10 ⁻⁵	0.292	2.53×10 ⁻²	4.18×10 ⁻³	6.59×10⁻⁵	0.261
TUT Dup.	1.43×10 ⁻³	4.59×10 ⁻⁴	7.59×10 ⁻⁵		2.69×10 ⁻²	4.48×10 ⁻³	6.14×10 ⁻⁵	

^a [RN] = Radionuclide concentration ^b Total propagated uncertainty ^c Minimum detectable concentration

^d Relative Error Ratio

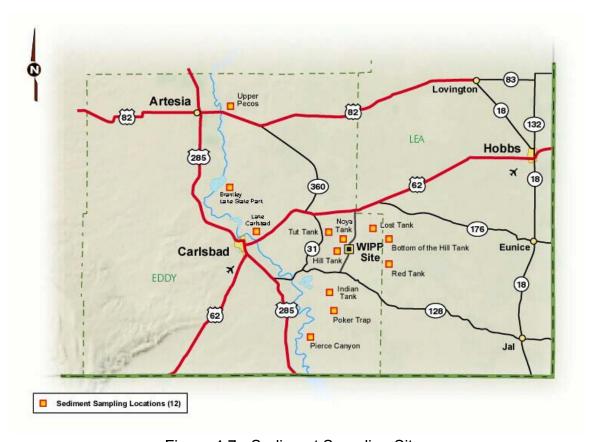


Figure 4.7 - Sediment Sampling Sites

4.8 Biota

4.8.1 Sample Collection

The concentration of radionuclides in plants is an important factor in estimating the intake of individual radionuclides by humans through ingestion. Therefore, rangeland vegetation samples were collected from the same six locations from where the soil samples were collected (Figure 4.6). The vegetation samples were chopped into 2.5-5-cm (1-2-in)-pieces, mixed together well, air dried at room temperature, and sent to the contract laboratory for analysis. Also collected were muscle tissues from two road-killed deer and one quail, both species commonly consumed by humans. Fish is also consumed in large amounts; therefore, fish samples from PCN, PEC, and BRA (three different locations on the Pecos River) were collected. The muscle tissues from the deer, quail, rabbit, and fish were sent to the laboratory for analysis.

4.8.2 Sample Preparation

Weighed aliquots were taken from the bulk of the chopped vegetation samples and animal tissue samples from each location. The aliquots were transferred into separate containers and dried at 100°C (212°F). Gamma spectrometric determinations of ⁴⁰K, ⁶⁰Co, and ¹³⁷Cs were performed directly from these aliquots. The samples were then dry-ashed, followed by wet-ashing and dissolution in 8M nitric acid. Aliquots from the dissolved samples were taken for the determinations of ⁹⁰Sr, ²³³⁺²³⁴U, ²³⁵U, ²³⁸U, ²³⁸Pu, ²³⁹⁺²⁴⁰Pu, and ²⁴¹Am.

4.8.3 Results and Discussions

Vegetation

Uranium-234 was detected in all vegetation samples; because of its naturally low concentration, ^{235}U was not detected in any vegetation sample (Table 4.19). Concentrations of ^{234}U ranged from $4.18\times10^{-4}\pm3.34\times10^{-4}$ Bq/g (1.13×10 $^{-2}\pm9.03\times10^{-3}$ pCi/g) at WSS to $8.21\times10^{-4}\pm2.75\times10^{-4}$ Bq/g (2.22×10 $^{-2}\pm7.43\times10^{-2}$ pCi/g) at WEE. Uranium-238 was detected all but one sample, WSS and varied between $2.54\times10^{-4}\pm1.32\times10^{-4}$ Bq/g (6.86×10 $^{-3}\pm3.57\times10^{-3}$ pCi/g) at MLR to $5.03\times10^{-4}\pm2.05\times10^{-4}$ Bq/g (1.36×10 $^{-2}\pm5.54\times10^{-3}$ pCi/g) at WEE. The concentration of ^{234}U and ^{238}U did not vary significantly between locations (ANOVA, ^{234}U p = 0.548, ^{238}U p = 0.527). The primary source for uranium in plant tissues is the soil, so this difference from the uranium results for soils may seem counterintuitive. However, uptake of radionuclides and contamination by resuspension are highly species dependent. Because of small-scale differences in soil type, shading, water availability, and other microenvironmental conditions, plants of the same species collected adjacent to one another will often have very different radionuclide concentrations.

Concentrations of ²³⁸Pu, ²³⁹⁺²⁴⁰Pu, and ²⁴¹Am were less than the minimum detectable concentrations in every vegetation sample (Table 4.19).

Potassium-40 was detected in every vegetation sample (Table 4.19), ranging from $1.64\times10^{-1}\pm2.75\times10^{-2}$ Bq/g $(4.43\times10^{0}\pm7.43\times10^{-1}$ pCi/g) at WEE to $4.29\times10^{-1}\pm6.48\times10^{-2}$ Bq/g $(1.16\times10^{1}\pm1.75\times10^{0}$ pCi/g) at SMR. The concentration of 40 K in vegetation was not significantly different between years (ANOVA, p = 0.348). Like uranium, the primary source for potassium in plant tissues is the soil, and this difference from the 40 K results for soil is probably due to the same factors. Cesium-137 was not detected in vegetation samples. Cobalt-60 was detected in sample location SEC. Strontium-90 was not detected in any of the vegetation samples.

Table 4.19 - Radionuclide Concentrations (Bq/g Wet Mass) in Vegetation Near the WIPP Site.

See Appendix B for the sampling locations.

	[RN] ^a	2×TPU ^b	MDC°	[RN]	2×TPU	MDC	[RN]	2×TPU	MDC
Location		²⁴¹ Am			²³⁸ Pu			²³⁹ Pu	
MLR	4.48×10 ⁻⁵	7.92×10 ⁻⁵	1.38×10 ⁻⁴	-1.59×10 ⁻⁵	5.51×10 ⁻⁵	1.47×10 ⁻⁴	-1.58×10 ⁻⁵	3.17×10 ⁻⁵	1.17×10 ⁻⁴
SEC	0.00×10 ⁰	0.00×10^{0}	1.53×10 ⁻⁴	1.31×10 ⁻⁵	6.92×10 ⁻⁵	1.40×10 ⁻⁴	0.00×10°	0.00×10^{0}	3.53×10 ⁻⁵
SMR	1.05×10 ⁻⁴	1.12×10 ⁻⁴	1.64×10 ⁻⁴	-6.48×10 ⁻⁵	2.25×10 ⁻⁴	5.99×10 ⁻⁴	0.00×10°	0.00×10^{0}	1.75×10 ⁻⁴
WEE	1.49×10 ⁻⁵	6.66×10 ⁻⁵	1.39×10 ⁻⁴	1.43×10 ⁻⁵	6.40×10 ⁻⁵	1.33×10 ⁻⁴	1.43×10 ⁻⁵	2.86×10 ⁻⁵	3.89×10 ⁻⁵
WFF	9.99×10 ⁻⁵	9.77×10 ⁻⁵	1.34×10 ⁻⁴	2.80×10 ⁻⁵	5.62×10 ⁻⁵	1.03×10 ⁻⁴	0.00×10°	0.00×10^{0}	1.03×10 ⁻⁴
WSS	3.22×10 ⁻⁵	6.48×10 ⁻⁵	1.19×10 ⁻⁴	4.55×10 ⁻⁵	6.81×10 ⁻⁵	1.11×10 ⁻⁴	0.00×10°	0.00×10^{0}	4.11×10 ⁻⁵
		²³⁴ U			²³⁵ U			²³⁸ U	
MLR	4.66×10 ⁻⁴	1.79×10 ⁻⁴	3.85×10 ⁻⁵	0.00×10 ⁰	0.00×10 ⁰	1.62×10 ⁻⁴	2.54×10 ⁻⁴	1.32×10 ⁻⁴	1.04×10 ⁻⁴
SEC	6.25×10 ⁻⁴	2.78×10 ⁻⁴	1.77×10 ⁻⁴	2.96×10 ⁻⁵	5.96×10 ⁻⁵	8.03×10 ⁻⁵	4.55×10 ⁻⁴	2.24×10 ⁻⁴	6.48×10 ⁻⁵
SMR	7.33×10 ⁻⁴	2.67×10 ⁻⁴	1.32×10 ⁻⁴	6.62×10 ⁻⁵	7.73×10 ⁻⁵	5.99×10⁻⁵	4.63×10 ⁻⁴	1.99×10 ⁻⁴	4.85×10⁻⁵
WEE	8.21×10 ⁻⁴	2.75×10 ⁻⁴	4.74×10 ⁻⁵	2.15×10 ⁻⁵	7.47×10 ⁻⁵	1.59×10 ⁻⁴	5.03×10 ⁻⁴	2.05×10 ⁻⁴	4.70×10 ⁻⁵
WFF	4.63×10 ⁻⁴	1.84×10 ⁻⁴	4.18×10 ⁻⁵	1.90×10 ⁻⁵	3.81×10 ⁻⁵	5.14×10 ⁻⁵	3.53×10 ⁻⁴	1.58×10 ⁻⁴	4.14×10 ⁻⁵
WSS	4.18×10 ⁻⁴	3.34×10 ⁻⁴	1.62×10 ⁻⁴	0.00×10°	0.00×10 ⁰	2.00×10 ⁻⁴	4.18×10 ⁻⁴	3.74×10 ⁻⁴	4.40×10 ⁻⁴
		¹³⁷ Cs			⁶⁰ Co				
MLR	6.77×10 ⁻⁴	1.15×10 ⁻³	1.39×10 ⁻³	1.19×10 ⁻³	1.54×10 ⁻³	1.87×10⁻³			
SEC	1.18×10 ⁻³	2.73×10 ⁻³	3.07×10 ⁻³	4.44×10 ⁻³	2.72×10 ⁻³	3.08×10 ⁻³			
SMR	6.14×10 ⁻⁴	2.64×10 ⁻³	2.96×10 ⁻³	1.68×10 ⁻³	2.71×10 ⁻³	3.01×10 ⁻³			
WEE	-2.75×10 ⁻⁶	1.17×10 ⁻³	1.37×10 ⁻³	3.14×10 ⁻⁵	1.53×10 ⁻³	1.78×10 ⁻³			
WFF	1.69×10 ⁻⁴	1.18×10 ⁻³	1.40×10 ⁻³	6.73×10 ⁻⁴	1.59×10 ⁻³	1.88×10 ⁻³			
WSS	9.66×10 ⁻⁴	1.68×10 ⁻³	1.95×10 ⁻³	9.73×10 ⁻⁴	1.86×10 ⁻³	2.29×10 ⁻³			
		90Sr			⁴⁰ K				
MLR	3.28×10 ⁻⁴	4.39×10 ⁻³	6.60×10 ⁻³	2.16E-01	3.53×10 ⁻²	1.78×10 ⁻²			
SEC	1.94×10 ⁻³	4.17×10 ⁻³	6.15×10 ⁻³	4.18E-01	6.36×10 ⁻²	3.05×10 ⁻²			
SMR	3.75×10 ⁻³	4.11×10 ⁻³	5.96×10 ⁻³	4.29E-01	6.48×10 ⁻²	2.76×10 ⁻²			
WEE	-4.23×10 ⁻⁴	4.42×10 ⁻³	6.69×10 ⁻³	1.64E-01	2.75×10 ⁻²	1.36×10 ⁻²			
WFF		4.20×10 ⁻³		2.93E-01	4.55×10 ⁻²	1.72×10 ⁻²			
WSS	1.47×10 ⁻³	4.41×10 ⁻³	6.54×10 ⁻³	2.56E-01	4.22×10 ⁻²	2.00×10 ⁻²			

^a [RN] = Radionuclide concentration

A duplicate analysis of the vegetation sample from WEE was performed for all the radionuclides of interest (Table 4.20). Concentrations of ²³⁴U, ²³⁸U, and ⁴⁰K were above detection limits in the duplicate sample. Relative Error Ratio values exceeded one for ²³⁴U, ²³⁸U, and ⁴⁰K, indicating a nonhomogenous sample.

^b Total propagated uncertainty

^c Minimum detectable concentration

Animals

Of the radionuclides of interest, ²³⁴U, ²³⁸U, and ⁴⁰K were detected in deer, rabbit, and quail tissue (Table 4.21). The mean concentrations did not differ from year 2000. These results can be used only as a gross indication of uptakes, as the sample sizes are too small to provide a robust analysis.

Table 4.20 - Results of Duplicate Vegetation Sample Analysis. Units are Bq/g. See Appendix B for the sampling locations.

	[RN] ^a	2×TPU ^b	MDC°	RER⁴	[RN]	2×TPU	MDC	RER			
Location		²³⁴ U				⁴⁰ K	⁴⁰ K				
WEE	8.21×10 ⁻⁴	2.75×10 ⁻⁴	4.74×10 ⁻⁵	1.839	1.64×10 ⁻¹	2.75×10 ⁻²	1.36×10 ⁻²	3.778			
WEE Dup.	2.43×10 ⁻⁴	1.52×10 ⁻⁴	1.61×10 ⁻⁴		4.22×10 ⁻¹	6.25×10 ⁻²	1.79×10 ⁻²				
		²³⁸ U									
WEE	5.03×10 ⁻⁴	2.05×10 ⁻⁴	4.70×10 ⁻⁵	1.14							
WEE Dup.	2.25×10 ⁻⁴	1.30×10 ⁻⁴	4.70×10 ⁻⁵								

a [RN] = Radionuclide concentration

Table 4.21 - Radionuclide Concentrations (Bq/g Wet Mass) in Deer and Quail Near the WIPP Site

	[RN] ^a	2×TPU ^b	MDC°	[RN]	2×TPU	MDC	[RN]	2×TPU	MDC		
Sample Type		²⁴¹ Am			²³⁸ Pu			²³⁹ Pu			
Deerd	3.14×10 ⁻⁷	1.42×10 ⁻⁶	1.32×10 ⁻⁶	2.74×10 ⁻⁷	1.04×10 ⁻⁶	1.59×10 ⁻⁶	-2.05×10 ⁻⁸	6.95×10 ⁻⁸	1.56×10 ⁻⁶		
Quaile	1.23×10 ⁻⁶	3.40×10 ⁻⁶	3.03×10 ⁻⁵	-1.01×10 ⁻⁴	2.79×10 ⁻⁴	2.87×10 ⁻⁵	0.00×10^{0}	0.00×10^{0}	8.35×10 ⁻⁶		
Rabbit ^f	1.84×10 ⁻⁶	2.13×10 ⁻⁶	1.65×10 ⁻⁶	0.00×10°	0.00×10°	1.67×10 ⁻⁶	-6.18×10 ⁻⁷	1.24×10 ⁻⁶	4.55×10 ⁻⁶		
		²³⁴ U			²³⁵ U			²³⁸ U			
Deer	1.07×10 ⁻⁵	2.62×10 ⁻⁵	5.51×10 ⁻⁷	1.44×10 ⁻⁷	3.74×10 ⁻⁷	6.80×10 ⁻⁷	7.92×10 ⁻⁶	1.93×10 ⁻⁵	5.49×10 ⁻⁷		
Quail	1.07×10 ⁻³	2.93×10 ⁻³	8.93×10 ⁻⁶	3.69×10 ⁻⁵	1.06×10 ⁻⁴	1.41×10 ⁻⁵	1.20×10 ⁻³	3.32×10 ⁻³	2.18×10 ⁻⁵		
Rabbit	4.81×10 ⁻⁵	1.31×10 ⁻⁵	1.61×10 ⁻⁶	5.11×10 ⁻⁶	3.96×10 ⁻⁶	1.98×10 ⁻⁷	3.89×10 ⁻⁵	1.14×10 ⁻⁵	1.60×10 ⁻⁶		
		¹³⁷ Cs		⁶⁰ Co							
Deer	-1.76×10 ⁻⁴	7.60×10 ⁻⁴	3.49×10 ⁻⁴	2.06×10 ⁻⁴	2.43×10 ⁻⁴	3.85×10 ⁻⁴	•				
Quail	2.09×10 ⁻⁴	2.02×10 ⁻³	3.95×10 ⁻³	1.77×10 ⁻³	2.04×10 ⁻³	5.16×10 ⁻³					
Rabbit	-1.59×10 ⁻⁴	5.59×10 ⁻⁴	6.40×10 ⁻⁴	-4.85×10 ⁻⁴	7.70×10 ⁻⁴	8.21×10 ⁻⁴					
		90Sr			⁴⁰ K		•				
Deer	-9.77×10 ⁻⁶	3.03×10 ⁻⁵	7.29×10 ⁻⁵	9.34×10 ⁻²	3.85×10 ⁻²	3.22×10 ⁻³	•				
Quail	1.55×10 ⁻³	2.16×10 ⁻³	1.54×10 ⁻³	1.41×10 ⁻¹	3.66×10 ⁻¹	4.23×10 ⁻²					
Rabbit	-9.95×10 ⁻⁵	1.38×10 ⁻⁴	2.33×10 ⁻⁴	1.07×10 ⁻¹	1.74×10 ⁻²	9.29×10 ⁻³					
a [DN] =	Radionuclio	de concentr	ation								

^a [RN] = Radionuclide concentration

Uranium-234 and ²³⁸U were detected in all the fish samples. Uranium-235 was detected in 67 percent of the fish samples (Table 4.22). Neither plutonium or ²⁴¹Am isotope was detected in fish.

^b Total propagated uncertainty

^c Minimum detectable concentration

d Relative Error Ratio

^b Total propagated uncertainty

^c Minimum detectable concentration

^d Mean of three samples collected near WIPP. TPU represents the standard deviation of the mean.

^e Mean of two samples collected near WIPP. TPU represents the standard deviation of the mean.

f Single Sample

Cesium-137, 60 Co, and 90 Sr were not detected in any of the fish samples (Table 4.22). Potassium-40 was detected in all fish (Table 4.22). It was lowest in the sample from BRA ($5.66\times10^{-2}\pm9.40\times10^{-3}$ Bq/g [$1.53\times10^{0}\pm2.54\times10^{-1}$ pCi/g]), and highest in the sample from PCN ($1.03\times10^{-1}\pm1.55\times10^{-2}$ Bq/g [$2.78\times10^{-1}\pm4.19\times10^{-1}$ pCi/g]).

Table 4.22 - Radionuclide Concentrations (Bq/g Wet Mass) in Fish Near the WIPP Site.

See Appendix B for the sampling locations.

	[RN] ^a	2×TPU ^b	MDC°	[RN]	2×TPU	MDC	[RN]	2×TPU	MDC
Location		²⁴¹ Am			²³⁸ Pu			²³⁹ Pu	
BRA	1.65×10 ⁻⁶	1.93×10 ⁻⁶	1.49×10 ⁻⁶	9.95×10 ⁻⁷	1.42×10 ⁻⁶	1.35×10 ⁻⁶	4.96×10 ⁻⁷	9.99×10 ⁻⁷	1.35×10 ⁻⁶
BRA	1.03×10 ⁻⁶	1.20×10 ⁻⁶	9.29×10 ⁻⁷	1.10×10 ⁻⁶	1.57×10 ⁻⁶	1.49×10 ⁻⁶	0.00×10^{0}	0.00×10 ⁰	1.49×10 ⁻⁶
PCN	6.48×10 ⁻⁷	1.30×10 ⁻⁶	1.75×10 ⁻⁶	5.96×10 ⁻⁷	1.20×10 ⁻⁶	1.61×10 ⁻⁶	5.92×10 ⁻⁷	1.19×10 ⁻⁶	1.61×10 ⁻⁶
PCN	1.28×10 ⁻⁶	1.82×10 ⁻⁶	1.73×10⁻ ⁶	1.26×10 ⁻⁶	1.47×10 ⁻⁶	1.14×10 ⁻⁶	0.00×10^{0}	0.00×10 ⁰	3.09×10 ⁻⁶
PEC	7.03×10 ⁻⁷	2.44×10 ⁻⁶	5.18×10 ⁻⁶	3.50×10 ⁻⁶	5.25×10 ⁻⁶	8.58×10 ⁻⁶	0.00×10°	0.00×10°	8.58×10 ⁻⁶
PEC	0.00×10°	0.00×10^{0}	4.92×10 ⁻⁶	-1.22×10 ⁻⁶	1.74×10 ⁻⁶	5.67×10 ⁻⁶	0.00×10°	0.00×10 ⁰	4.48×10 ⁻⁶
		²³⁴ U			²³⁵ U			²³⁸ U	
BRA	8.70×10 ⁻⁵	1.68×10 ⁻⁵	8.29×10 ⁻⁷	3.02×10 ⁻⁶	2.18×10 ⁻⁶	1.02×10 ⁻⁶	3.77×10 ⁻⁵	8.92×10 ⁻⁶	8.25×10 ⁻⁷
BRA	5.48×10 ⁻⁵	1.08×10 ⁻⁵	5.88×10 ⁻⁷	1.07×10 ⁻⁶	1.08×10 ⁻⁶	7.25×10 ⁻⁷	3.26×10 ⁻⁵	7.29×10 ⁻⁶	5.85×10 ⁻⁷
PCN	1.03×10 ⁻⁴	2.19×10 ⁻⁵	4.33×10 ⁻⁶	4.00×10 ⁻⁶	3.10×10 ⁻⁶	1.55×10 ⁻⁶	5.37×10 ⁻⁵	1.32×10 ⁻⁵	1.25×10 ⁻⁶
PCN	1.58×10 ⁻⁴	2.99×10 ⁻⁵	9.88×10 ⁻⁷	3.59×10 ⁻⁶	2.60×10 ⁻⁶	1.22×10 ⁻⁶	8.51×10 ⁻⁵	1.79×10⁻⁵	2.67×10 ⁻⁶
PEC	1.32×10 ⁻⁴	2.70×10 ⁻⁵	1.65×10⁻ ⁶	2.25×10 ⁻⁶	2.62×10 ⁻⁶	2.04×10 ⁻⁶	6.73×10 ⁻⁵	1.64×10 ⁻⁵	1.64×10 ⁻⁶
PEC	1.96×10 ⁻⁴	3.63×10 ⁻⁵	1.11×10 ⁻⁶	5.07×10 ⁻⁶	3.31×10 ⁻⁶	1.37×10 ⁻⁶	9.62×10 ⁻⁵	2.00×10 ⁻⁵	3.81×10 ⁻⁶
		¹³⁷ Cs			⁶⁰ Co				
BRA	9.99×10 ⁻⁵	6.55×10 ⁻⁴	7.62×10 ⁻⁴	-1.65×10 ⁻⁴	7.33×10 ⁻⁴	7.81×10 ⁻⁴			
BRA	-4.29×10 ⁻⁴	5.33×10 ⁻⁴	5.59×10 ⁻⁴	1.66×10⁻⁴	5.00×10 ⁻⁴	5.77×10 ⁻⁴			
PCN	3.92×10 ⁻⁴	1.01×10 ⁻³	1.18×10 ⁻³	-5.07×10 ⁻⁴	1.13×10 ⁻³	1.18×10 ⁻³			
PCN	-3.45×10 ⁻⁴	5.44×10 ⁻⁴	5.77×10 ⁻⁴	2.06×10 ⁻⁴	5.25×10 ⁻⁴	6.07×10 ⁻⁴			
2.37×10 ⁻³									
PEC		1.39×10 ⁻³	1.63×10 ⁻³	3.89×10⁻⁵	1.49×10 ⁻³	1.62×10 ⁻³			
PEC	2.12×10 ⁻⁴		6.03×10 ⁻⁴	-2.85×10 ⁻⁴	6.29×10 ⁻⁴	6.85×10 ⁻⁴			
		90Sr			⁴⁰ K				
BRA	1.45×10 ⁻⁵		1.54×10 ⁻⁴	9.51×10 ⁻²	1.69×10 ⁻²	1.51×10 ⁻²			
BRA	3.35×10 ⁻⁵		1.26×10 ⁻⁴	5.66×10 ⁻²	9.40×10 ⁻³	6.29×10 ⁻³			
PCN	2.20×10 ⁻⁴	1.69×10 ⁻⁴	2.60×10 ⁻⁴	8.88×10 ⁻²	1.98×10 ⁻²	2.29×10 ⁻²			
PCN	8.55×10 ⁻⁵		1.13×10 ⁻⁴	1.03×10 ⁻¹	1.55×10 ⁻²	6.66×10 ⁻³			
PEC	7.03×10 ⁻⁵	1.76×10 ⁻⁴	2.83×10 ⁻⁴	9.14×10 ⁻²	2.46×10 ⁻²	3.18×10 ⁻²			
PEC	-2.73×10 ⁻⁵	1.02×10 ⁻⁴	1.70×10 ⁻⁴	8.81×10 ⁻²	1.47×10 ⁻²	8.62×10 ⁻³			

^a [RN] = Radionuclide concentration

Neither ²⁴¹Am nor any plutonium isotopes were detected in fish.

4.9 **Summary and Conclusion**

The Environmental Monitoring Program collected samples of air particulates, soil, sediment, groundwater, surface water, and biota and analyzed them for radionuclides considered to be indicators of potential contamination from the WIPP facility, as well as other radionuclides of potential interest. Measured concentrations were examined for evidence of WIPP-related contamination, such as higher concentrations of TRU

^b Total propagated uncertainty

^c Minimum detectable concentration

radionuclides after 1998, or higher concentrations in downwind or down gradient directions. Radionuclide concentrations observed were highly variable in space and time and between media. However, no time or space relationships related to WIPP were observed, and concentrations were consistent with background levels. In no case, could environmental concentrations be attributed to WIPP releases. In addition, no events occurred at WIPP, which would lead one to suspect that a release had occurred.

